Attorney's Docket No.: 21128-002US1 / B0247WO/US

Applicant: David Klatzmann et al.

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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

- 1 (Currently Amended) Method for obtaining, preparing or producing human suppressor T lymphocytes [[(]]and/or the precursors thereof[[)]], comprising a step of selection, separation or isolation in vitro or ex vivo of human T lymphocytes expressing the THY-1 molecule.
 - 2. (Original) Method according to claim 1, comprising:
 - (a) obtaining a cell population of human origin comprising T lymphocytes, and
 - (b) recovering T lymphocytes expressing the THY-1 antigen.
- 3. (Currently Amended) Method according to claim 1, characterized in that wherein the step (b) is preceded or followed by a step of amplification of T lymphocytes.
- 4. (Currently Amended) Method according to any one of the previous claims, characterized in that The method of claim 1 wherein the T lymphocytes expressing the THY-1 antigen are selected, separated, isolated or recovered by means of a ligand specific of THY-1.
- 5. (Currently Amended) Method according to claim 4, characterized in that wherein the specific ligand is an antibody specific of THY-1 or a fragment or derivative of said antibody having substantially the same antigenic specificity.
- 6. (Currently Amended) Method according to claim 5, characterized in that wherein the specific ligand is a monoclonal or polyclonal antibody specific of THY-1.
- 7. (Currently Amended) Method according to claim 5, characterized in that wherein the specific ligand is a polyfunctional, monocatenary or multimeric antibody, specific of THY-1.

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8. (Currently Amended) Method according to claim 4, characterized in that wherein the specific ligand is an aptamer.

- 9. (Currently Amended) Method according to any one of claims 4 to 8 claim 4, characterized in that wherein the ligand is immobilized on a support or placed in solution.
- 10. (Currently Amended) Method according to claim 9, characterized in that wherein the support is a column or a bead, preferably a magnetic bead.
- 11. (Currently Amended) Method according to any one of claims 4 to 10 claim 4, characterized in that wherein the ligand is labelled.
- 12. (Currently Amended) Method according to claim 11, characterized in that wherein the labelling is carried out by means of a fluorescent, radioactive, luminescent, phosphorescent, chemical or enzymatic detection label.
- 13. (Currently Amended) Method according to any one of the previous claims claim 1, characterized in that wherein the step of recovery, selection or isolation is carried out by flow cytometry, affinity chromatography, FACS, MACS or D/MACS.
- 14. (Currently Amended) Method according to any one of the previous claims claim 2, characterized in that wherein the cell population comes from a tissue selected in the group consisting of bone marrow, spleen, liver, thymus, blood which has or has not been previously enriched in T lymphocytes, umbilical cord blood, fetal, infant or adult peripheral blood, a tumor, a site of inflammation, a transplanted organ or a cell culture established with one or another of said tissues.

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15. (Original) Method for identifying and/or quantifying human suppressor T lymphocytes in a cell population, comprising exposing said cell population to a ligand specific of THY-1 and determining and/or quantifying the formation of a complex between the ligand and the cells, the formation of said complexes indicating the presence and/or the quantity of suppressor T lymphocytes in the cell population.

- 16. (Original) Method for producing a pharmaceutical composition, comprising:
- (a) obtaining a biological sample comprising human T lymphocytes,
- (b) selecting T lymphocytes expressing the THY-1 antigen in said biological sample, and
- (c) conditioning said T lymphocytes expressing the THY-1 antigen in a pharmaceutically acceptable adjuvant or medium.
 - 17. (Original) Method for producing a pharmaceutical composition, comprising:
 - (a) obtaining a biological sample comprising human T lymphocytes,
- (b) depleting T lymphocytes expressing the THY-1 antigen from said biological sample, and
- (c) conditioning said T lymphocytes not expressing the THY-1 antigen in a pharmaceutically acceptable adjuvant or medium.

Claims 18-28 (Canceled)